

**MAXIMUM STRENGTH ANTACID AND ANTIGAS- aluminum hydroxide,
magnesium hydroxide, dimethicone suspension
VALU MERCHANDISERS COMPANY**

BC max antacid

Active ingredients (in each 5 mL teaspoonful)

Aluminum hydroxide 400 mg (equivalent to dried gel, USP)

Magnesium hydroxide 400 mg

Simethicone 40mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- the symptoms referred to as gas

Warnings

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are taking a prescription drug.

Antacids may interact with certain prescription drugs.

Stop use and ask a doctor if symptoms last more than 2 weeks

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Directions

- shake well before use
- adults and children 12 years and older: take 2 to 4 teaspoonfuls between meals, at bedtime, or as directed by a doctor
- do not exceed 12 teaspoonfuls in a 24 hour period or use the maximum dosage for

- more than 2 weeks
- children under 12 years of age: ask a doctor

Other information

- **each 5 mL teaspoonful contains:** magnesium 165 mg, sodium 5 mg
- do not freeze
- store at room temperature tightly closed

Inactive ingredients

benzyl alcohol, butylparaben, carboxymethylcellulose sodium, flavor (contains alcohol), hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments?

1-800-540-3765


package Label

Best Choice[®] Compare to the active ingredients in
Maximum Strength Mylanta^{®*}

MAXIMUM STRENGTH

Antacid Anti-Gas

ALUMINA, MAGNESIA, AND SIMETHICONE ORAL SUSPENSION



Fast relief of:

- Acid indigestion
- Heartburn
- Sour stomach
- Pressure & bloating

ORIGINAL FLAVOR
 Alcohol: 0.5%

12 FL OZ (355 mL)

276-06122-2 REV0323

Drug Facts TAMPER EVIDENT: DO NOT USE IF THE BREAKAWAY BAND ON CAP IS BROKEN OR MISSING.

Active ingredients (in each 5 mL teaspoonful)	Purpose
Aluminum hydroxide (equivalent to dried gel, USP) 400 mg	Antacid
Magnesium hydroxide 400 mg	Antacid
Simethicone 40 mg	Antigas

Uses relieves • heartburn • sour stomach • acid indigestion • the symptoms referred to as gas

Warnings
 Ask a doctor before use if you have
 • kidney disease • a magnesium-restricted diet
 Ask a doctor or pharmacist before use if you are presently taking a prescription drug. Antacids may interact with certain prescription drugs.
 Stop use and ask a doctor if symptoms last for more than 2 weeks
 If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children.

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Questions or comments? 1-800-540-3765

*This product is not manufactured or distributed by P & G to McNeil Consumer Pharmaceuticals Company, distributor of Mylanta[®].
 PRODUCT DISTRIBUTED BY:
 ASSOCIATED WHOLESALE GROCERS, INC
 KANSAS CITY, KANSAS 66106
 For product information please visit:
 bestchoicebrand.com/20147

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 www.bestchoicebrand.com

MAXIMUM STRENGTH ANTACID AND ANTIGAS

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-619
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IU0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IU0)	ALUMINUM HYDROXIDE	400 mg in 5 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	400 mg in 5 mL
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	40 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-619-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	08/01/2017	

Labeler - VALU MERCHANDISERS COMPANY (868703513)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(63941-619) , pack(63941-619)

Revised: 10/2023

VALU MERCHANDISERS COMPANY